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CLAIM AMENDMENTS

1. (original) A compound of formula

$$\begin{bmatrix} R^4 \\ R^{3a} \\ R^3 \\ R^3 \end{bmatrix}$$

$$\begin{bmatrix} R^3 \\ R^3 \\ R^3 \\ R^4 \end{bmatrix}$$

$$\begin{bmatrix} R^4 \\ R^3 \\ R^3 \\ R^4 \end{bmatrix}$$

$$\begin{bmatrix} R^4 \\ R^3 \\ R^4 \end{bmatrix}$$

wherein

R¹ and R² are chosen from H, halogen, -OH, loweralkyl, -O-loweralkyl, -CN, -S-loweralkyl, amino, lower alkylamino, alkylsulfonyl, arylsulfonyl, acyl, a sugar, a glucuronide and a sugar carbamate;

R³ is chosen from H, -OH, fluoro and -O-loweralkyl;

R^{3a} is chosen from H and fluoro, or R^{3a} and R³ together are =O;

R⁴ is chosen from H, halogen, -OH, loweralkyl, -O-loweralkyl, -CN, -S-loweralkyl, amino, lower alkylamino, alkylsulfonyl, arylsulfonyl and acyl;

$$Q^a$$
 Q^b Q^b Q^c Q^c

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 Q^a , Q^b and Q^c are independently chosen from a direct bond, -O-, -S-, -NH-, -CH₂O-, -CH₂NH-, -OCH₂CONH-, -OCH₂COO-, -C(=O)-, -CONH-, -NHCO-, -O(C=O)-, -(C=O)O-, -NHCONH-, -OCONH- and -NHCOO-; n is 2 or 3; m is 0,1, 2 or 3 and m = n; and

A has a valency of n and is chosen from C₂ to C₂₀ hydrocarbon, substituted alkyl of 2 to 20 carbons, perfluoroalkyl of 2 to 20 carbons, substituted aryl, polyaryl of 3 to 20 aryl groups, substituted arylalkyl, oxaalkyl of four to fifty carbons, azaalkyl of four to fifty carbons, thiaalkyl of four to fifty carbons, a residue of an oligopeptide of two to twenty amino acids, a residue of a monosaccharide or of a polysaccharide of 2 to 100 saccharide residues; and, when Q^a and Q^b are -O(C=O)- or -NHCO-, A may additionally be methylene.

2. (original) A compound according to claim 1 wherein m = zero of formula:

$$R^{3a}$$
 R^3 R

3. (original) A compound according to claim 1 wherein m=n of formula:

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$$\mathbb{R}^{38}$$
 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3 \mathbb{R}^3

- 4. (currently amended) A compound according to any of claims 1-3 claim 1 wherein n is 3 and W is trivalent.
- 5. (original) A compound according to claim 4 wherein Q^a , Q^b and Q^c are independently chosen from -O-, -CH₂O-, -OCH₂CONH-, -OCH₂COO-, -(C=O)O-, and -NHCOO-; and A is a polysaccharide of 2 to 20 saccharide residues, a branched oxaalkyl of four to fifty carbons or a monoazaalkyl of four to ten carbons.
- 6. (original) A compound according to claim 4 wherein Q^a , Q^b and Q^c are independently chosen from -CH₂O-, -CH₂NH-, -OCH₂CONH-, -OCH₂COO-, -CONH-, -NHCO-, -O(C=O)-, -(C=O)O-, -NHCONH-, -OCONH- and -NHCOO-; and A is an oligopeptide.

7. (currently amended) A compound according to any of claims 1-3 claim 1 wherein n is 2 and W is divalent of formula:

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(original) A compound according to claim 7 wherein 8.

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 Q^a and Q^b are independently chosen from -O-, -CH2O-, -OCH2CONH-, -OCH2COO-, -(C=O)O-, and -NHCOO-; and

A is poly(oxyethylene) or a polysaccharide of 2 to 20 saccharide residues.

9. (original) A compound according to claim 7 wherein

$$Q^a$$
 and Q^b are independently chosen -CH₂O-, -CH₂NH-, -OCH₂CONH-, -OCH₂COO-, -CONH-, -NHCO-, -O(C=O)-, -(C=O)O-, -NHCONH-, -OCONH- and -NHCOO-; and

A is an oligopeptide.

10. (currently amended) A compound according to any of claims 1-9 claim 1 wherein R¹ and R² are chosen from H, halogen, -OH, and methoxy;

R³ is -OH; and

R⁴ is fluoro.

(currently amended) A compound according to any of claims 1-9 claim 1 wherein R¹ and R² are chosen from a sugar, a glucuronide and a sugar carbamate;
 R³ is -OH; and

R⁴ is fluoro.

12. (currently amended) A compound according to any of claims 1-10 claim 1 of formula:

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13. (original) A compound according to claim 12 wherein W is chosen from -OCH₂CH=CHCH₂O-,

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14. (original) A compound according to claim 1 chosen from the group consisting of

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and

- 15. (currently amended) A pharmaceutical formulation comprising a compound according to any of claims 1-14 claim 1 and a pharmaceutically acceptable carrier.
- 16. (original) A pharmaceutical formulation according to claim 15 additionally comprising an inhibitor of cholesterol biosynthesis.

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17. (currently amended) A method for treating a disorder of lipid metabolism comprising administering a to a mammal a therapeutically effective amount of a compound according to any of claims 1-14 claim 1.

- 18. (original) A method according to claim 17, wherein said disorder of lipid metabolism is hyperlipidemia.
- 19. (original) A method according to claim 17, wherein said disorder of lipid metabolism is arteriosclerosis.
- 20. (currently amended) A method for inhibiting the absorption of cholesterol from the intestine of a mammal, which comprises administering an effective cholesterol-absorption-inhibiting amount of a compound according to any of claims 1-14 claim 1 to the mammal.

21. - 30. (canceled)